



Acinetobacter baumannii Infection and Transmission in the Military Health Care System



James Mancuso
MAJ, MC
C, Epidemiology
USACHPPMEUR



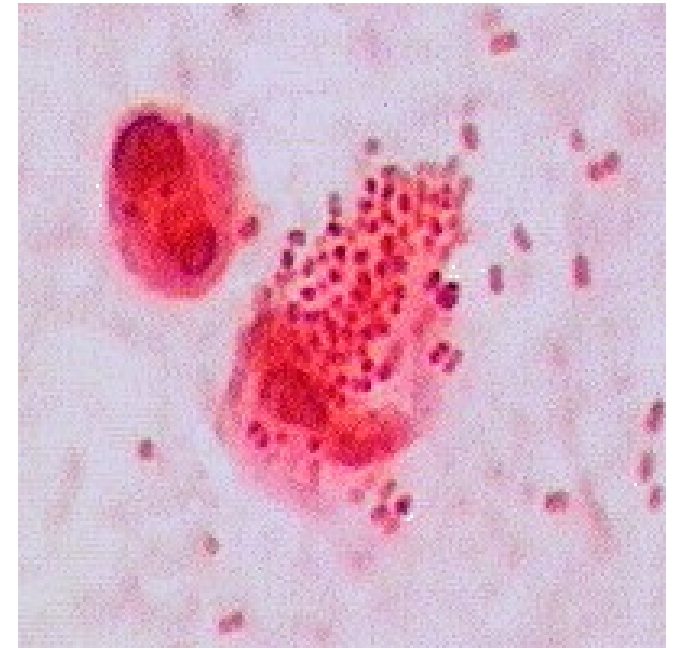
Outline

- Background
- *A. baumannii* transmission throughout MEDEVAC Sequence
 - Iraq
 - LRMC
 - WRAMC
- Infection control and nosocomial transmission
- Recommendations
- Future directions



Background

- Common nosocomial pathogen
 - Recognized pathogen in Vietnam war
 - *A. baumannii* seen almost exclusively in medical treatment facilities
 - VAP, Burn units, ICUs
- Infections, colonization or both noted
- Acinetobacter species are common skin contaminants, but not *A. baumannii*
- Difficult to eradicate
 - Very hard to eliminate (lives on surfaces for over 48 hrs)
 - Often is multi-drug resistant so acquisition is hard to treat
 - Persistence on human host is unknown (1-176 days in 1 recent abstract)

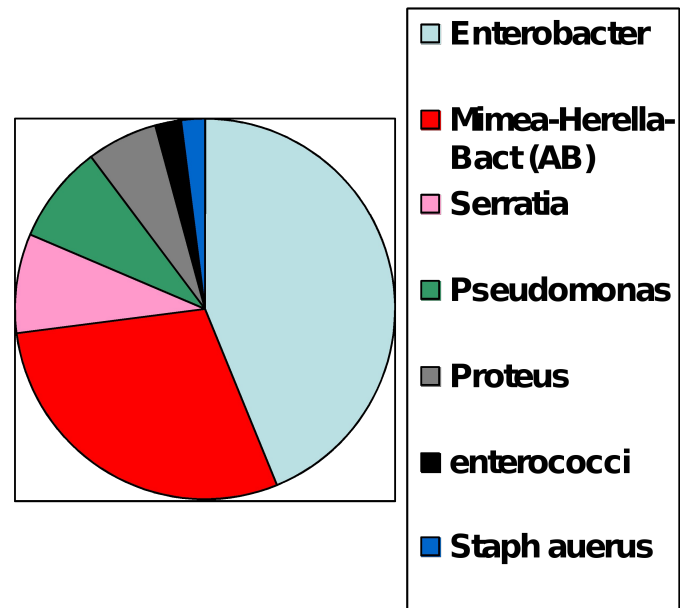




Acinetobacter in Vietnam

- Study of wound infections at DaNang hospital in 68-69
- 30 critically ill patients
- 12 patients with bacteremia
- *JAMA* 1972:219(8), 1044-7.

Bloodstream isolates by species





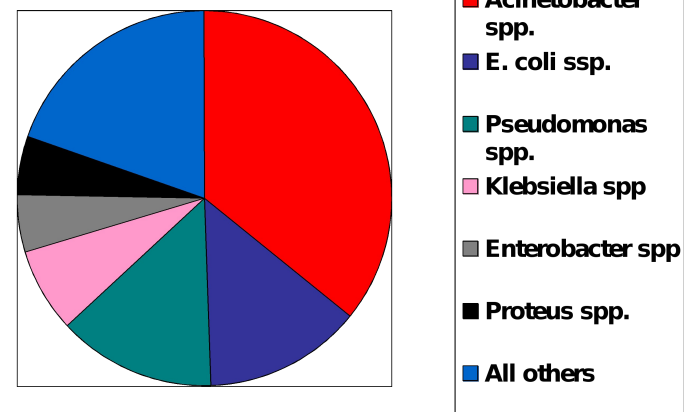
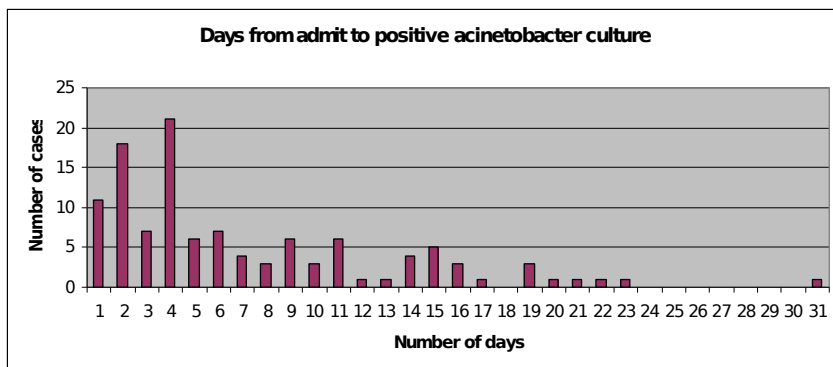
Timeline

- 19 Mar 2003: OIF begins
- 21 Mar 2003: First LRMC Isolates seen
- April 2003: USNS Comfort notes increase in isolates of highly resistant *A. baumannii*
- June 2003: LRMC begins active surveillance program
- Summer 2004: Several nosocomial deaths at WRAMC and LRMC
- Aug 2004: EPICON directed by TSG
- Nov 2004: MMWR article, other publications



USNS Comfort 2003

- 23% Infected or colonized with MDR *Acinetobacter* spp.
- Positive cultures early in hospital course
- 80% of patients were Iraqis



Source: LCDR Kyle Peterson, NNMC



Acinetobacter Theories

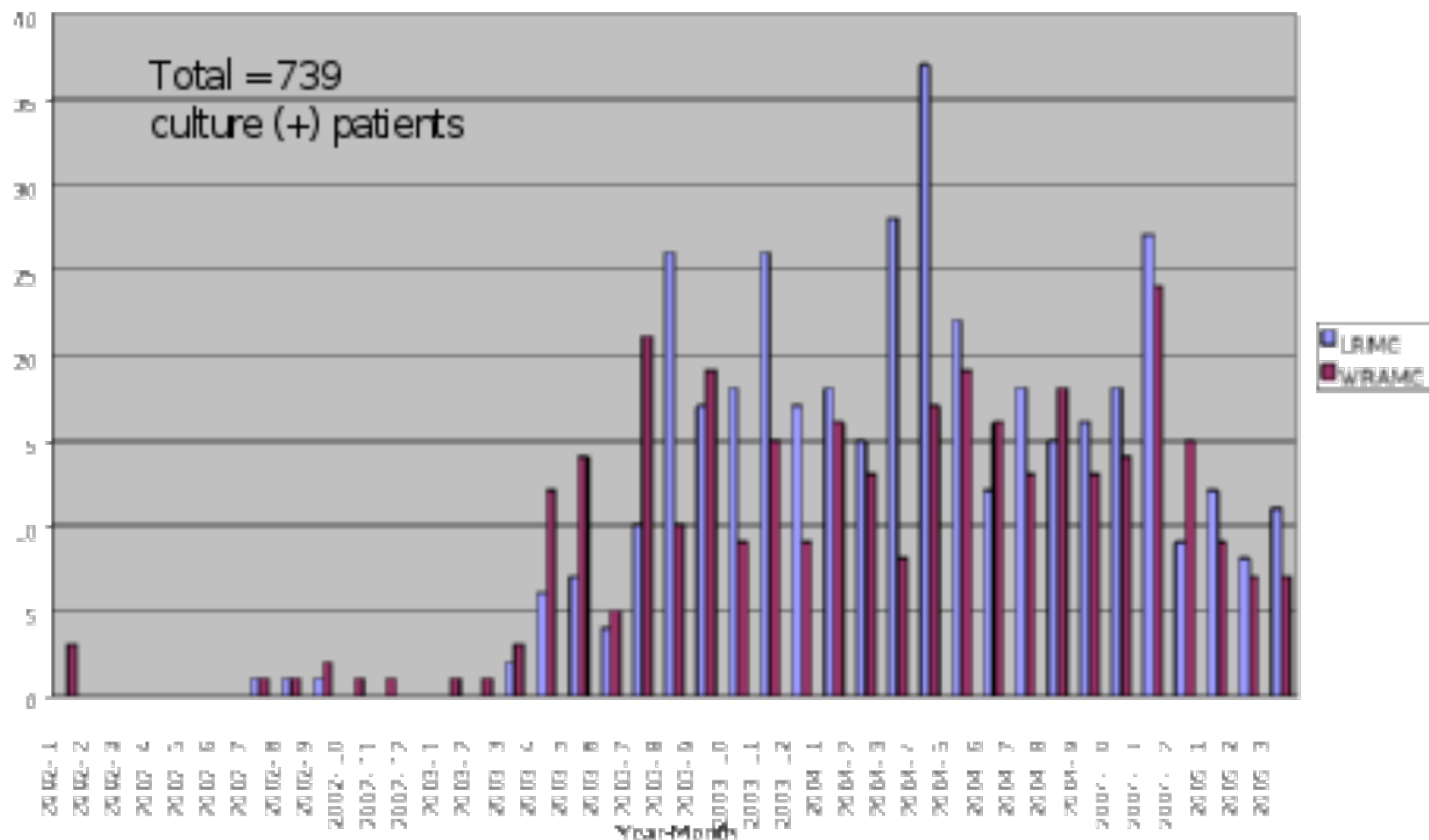
- Dirt in Iraq
- Propensity for colonization in the tropics
- Perioperative Ancef
- Cross-contamination of US troops by Iraqis
- Difficulty in eradication
 - Normal hospital setting difficult
 - Translation to MEDEVAC chain with multiple moving parts



Counts of First Positive *A. baumannii* Culture per Infected or Colonized Patient Walter Reed Army Medical Center & Landstuhl Regional Medical Center

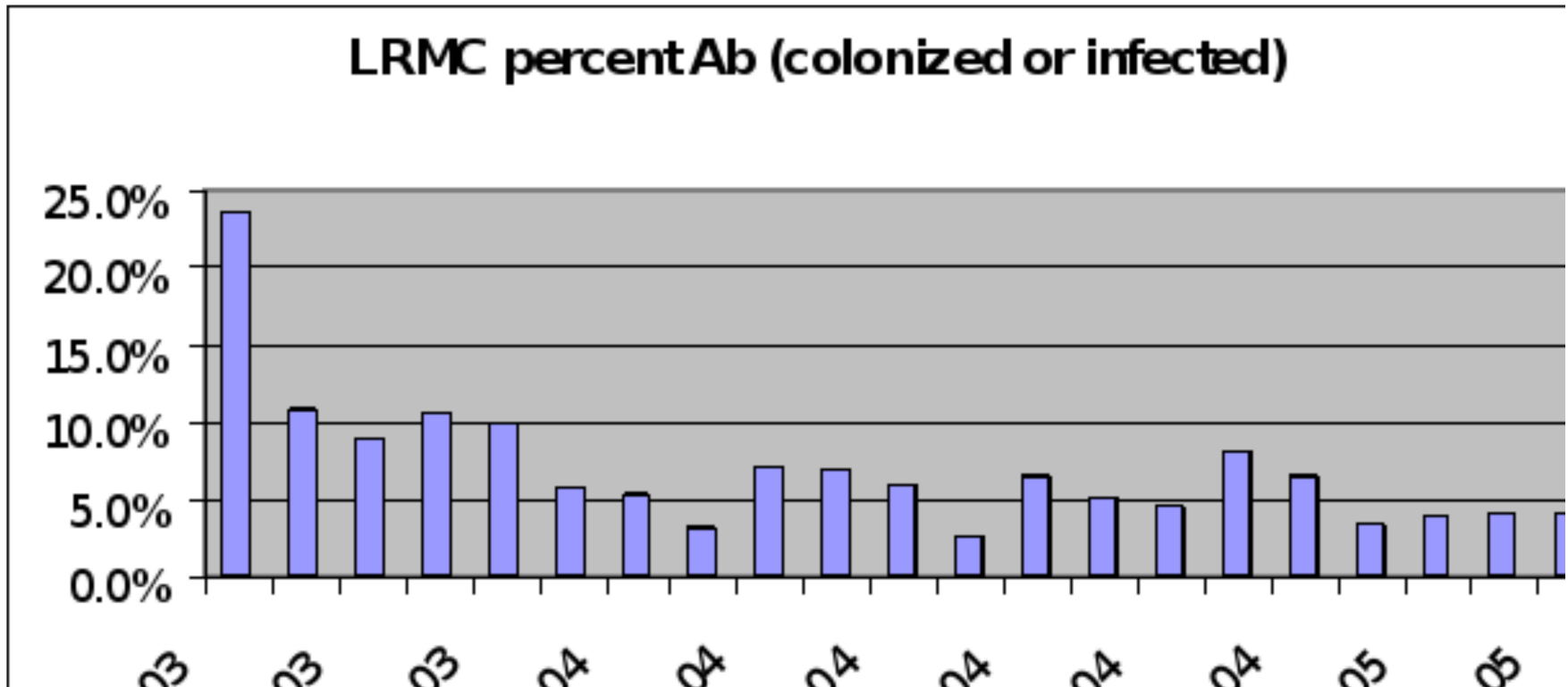
January 2002 - March 2005

BY HOSPITAL; ANY SPECIMEN TYPE





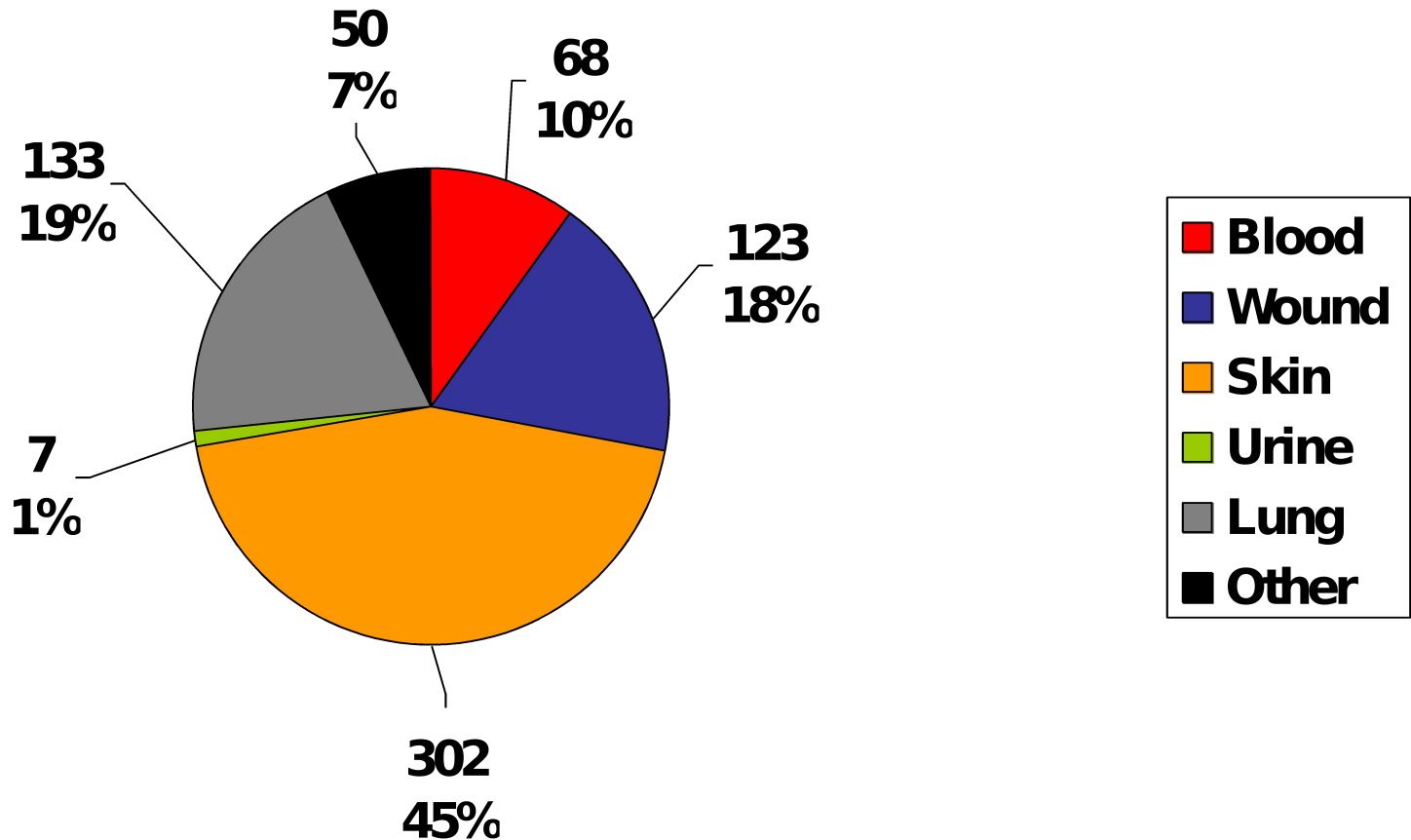
LRMC *A. baumannii* Infection Control Surveillance



Source: Greg Deye and Abel Trevino, LRMC



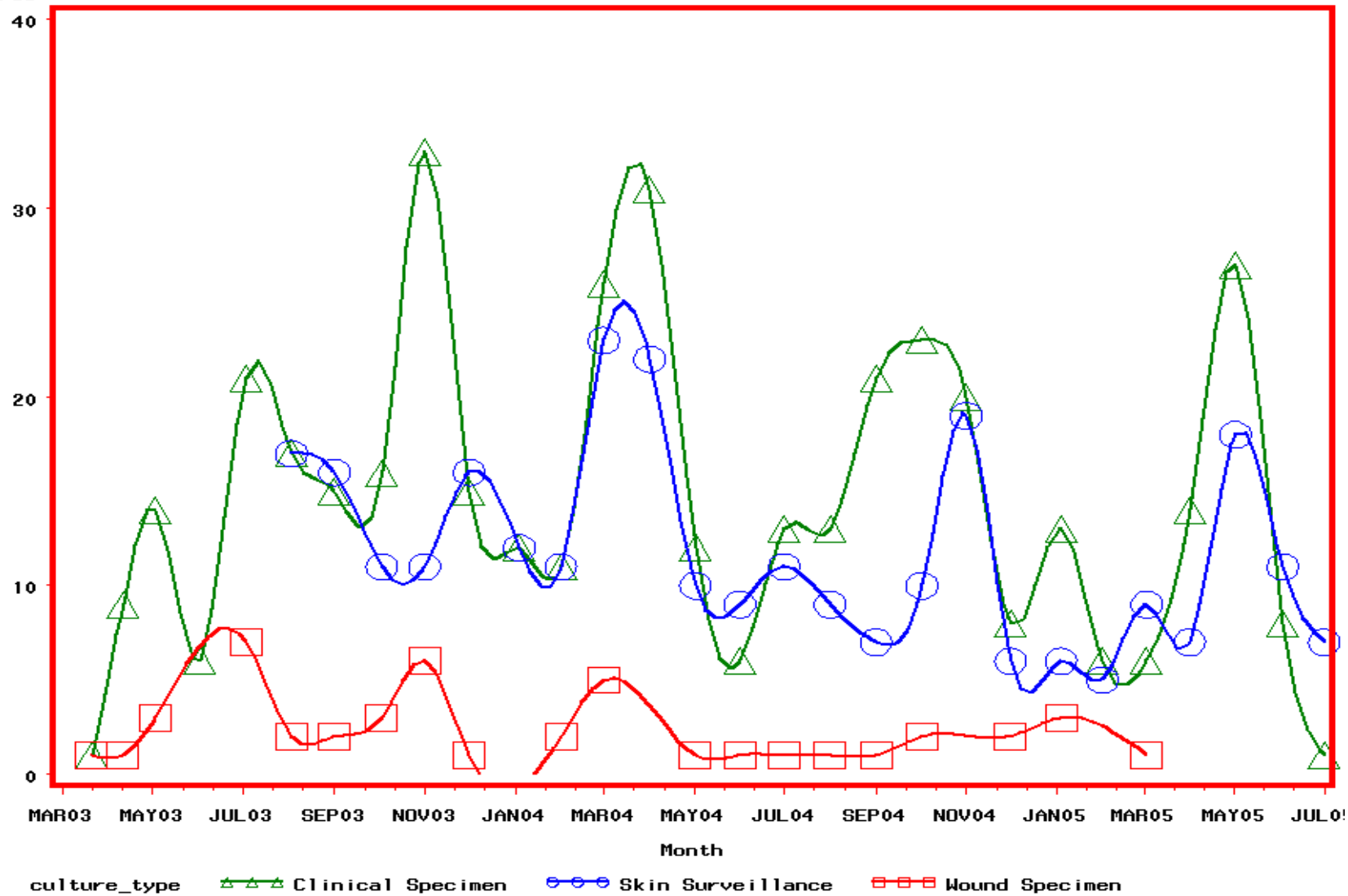
Sites of Acinetobacter Cultures, LRMC, January 2002 to April 2005



N=683

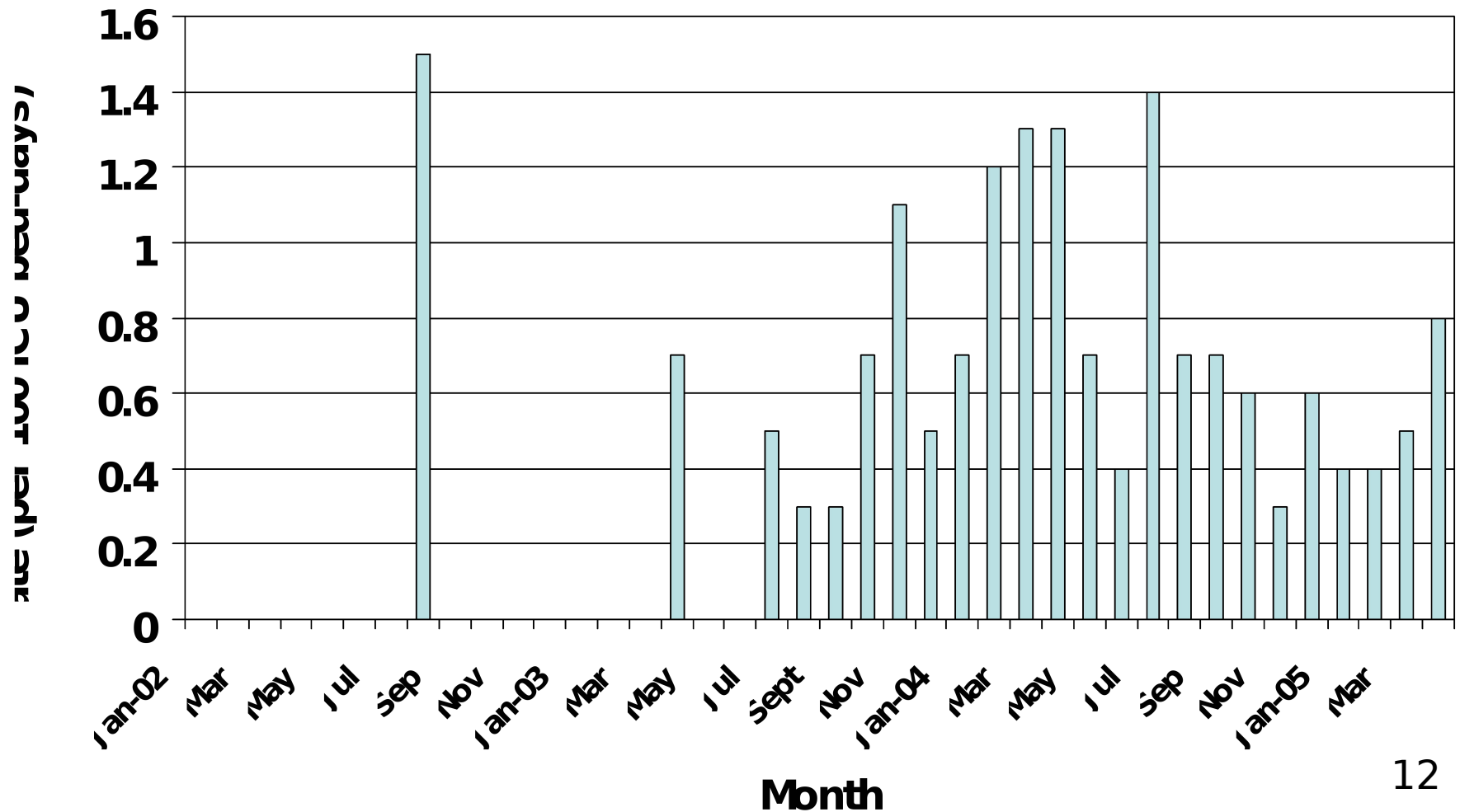
Acinetobacter baumannii Surveillance and Clinical Cultures, Landstuhl Regional Medical Center, 2003—2005

Positive cultures



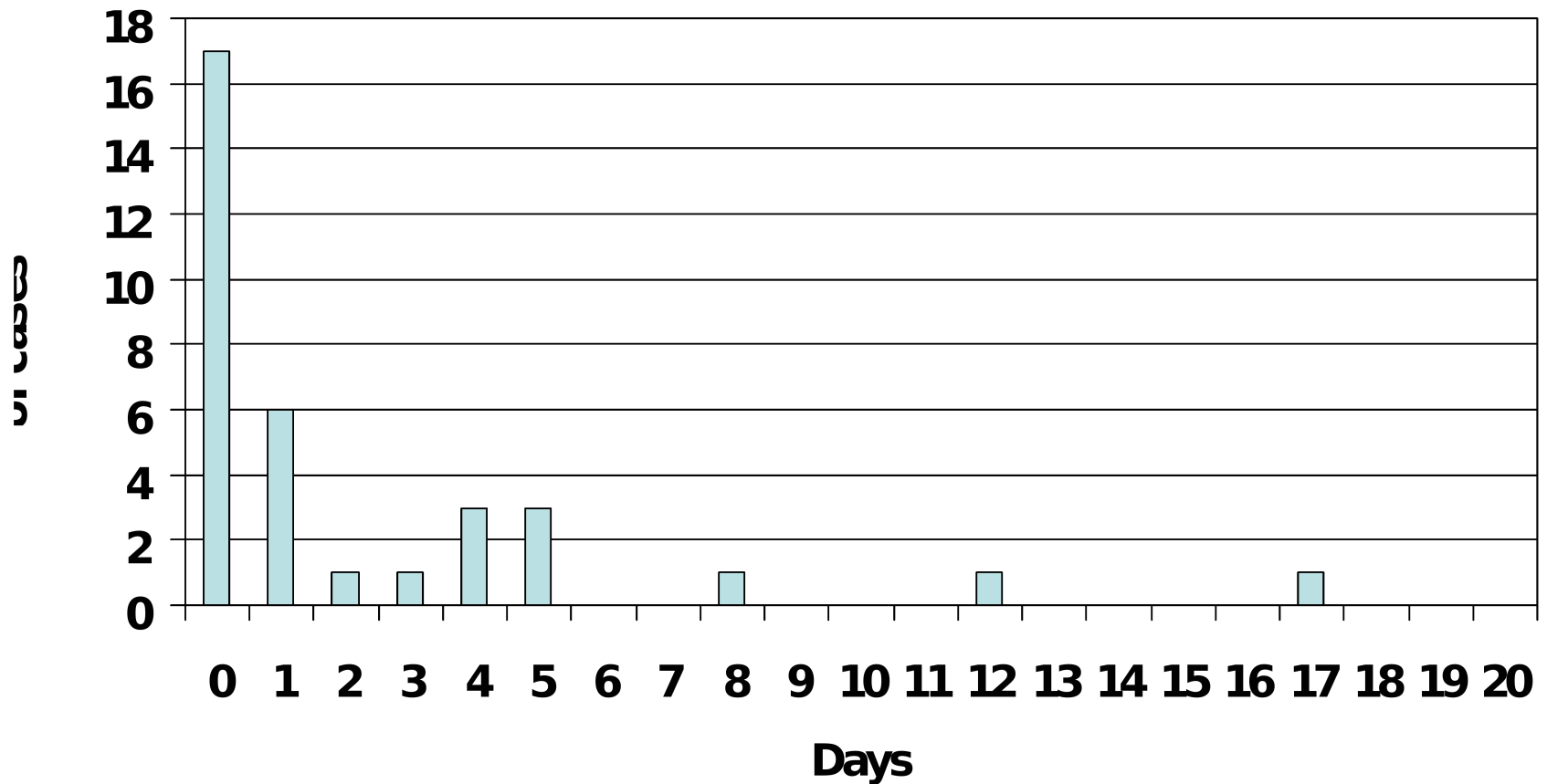


Rate of Acinetobacter Bloodstream Infections, 2002- 2005



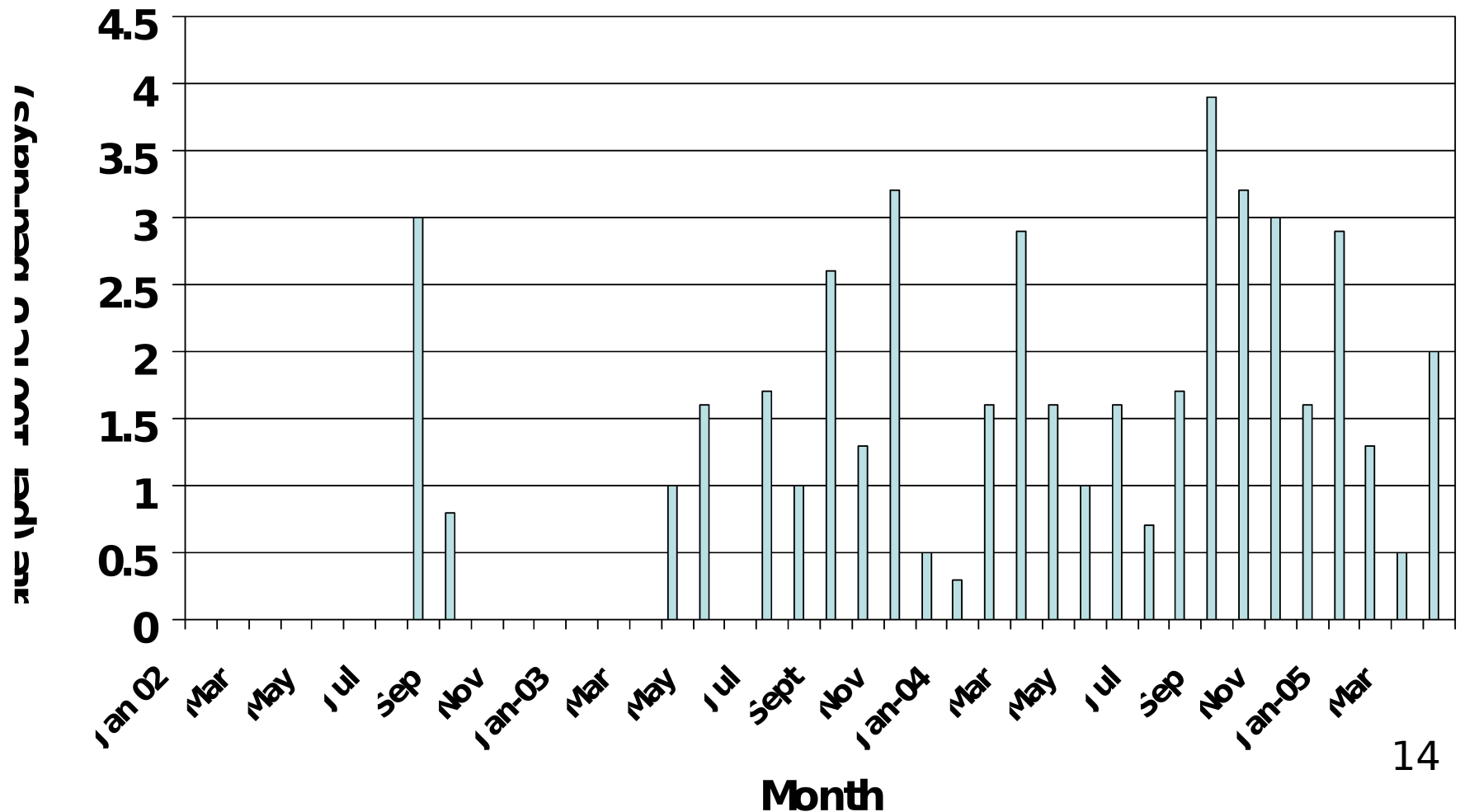


Time from Admission to Collection of Positive Blood Culture



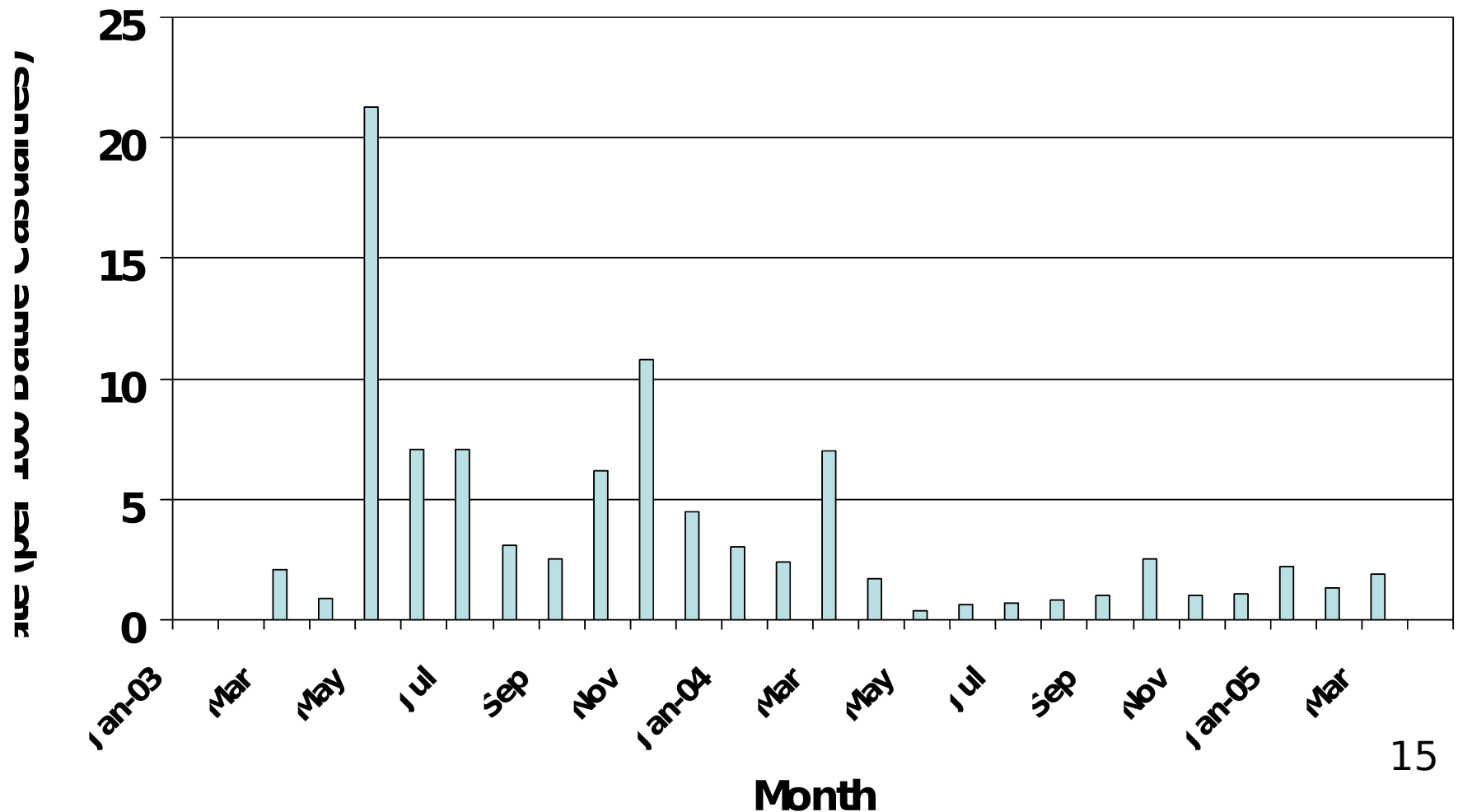


Rate of Respiratory Cultures with Acinetobacter, 2002-2005



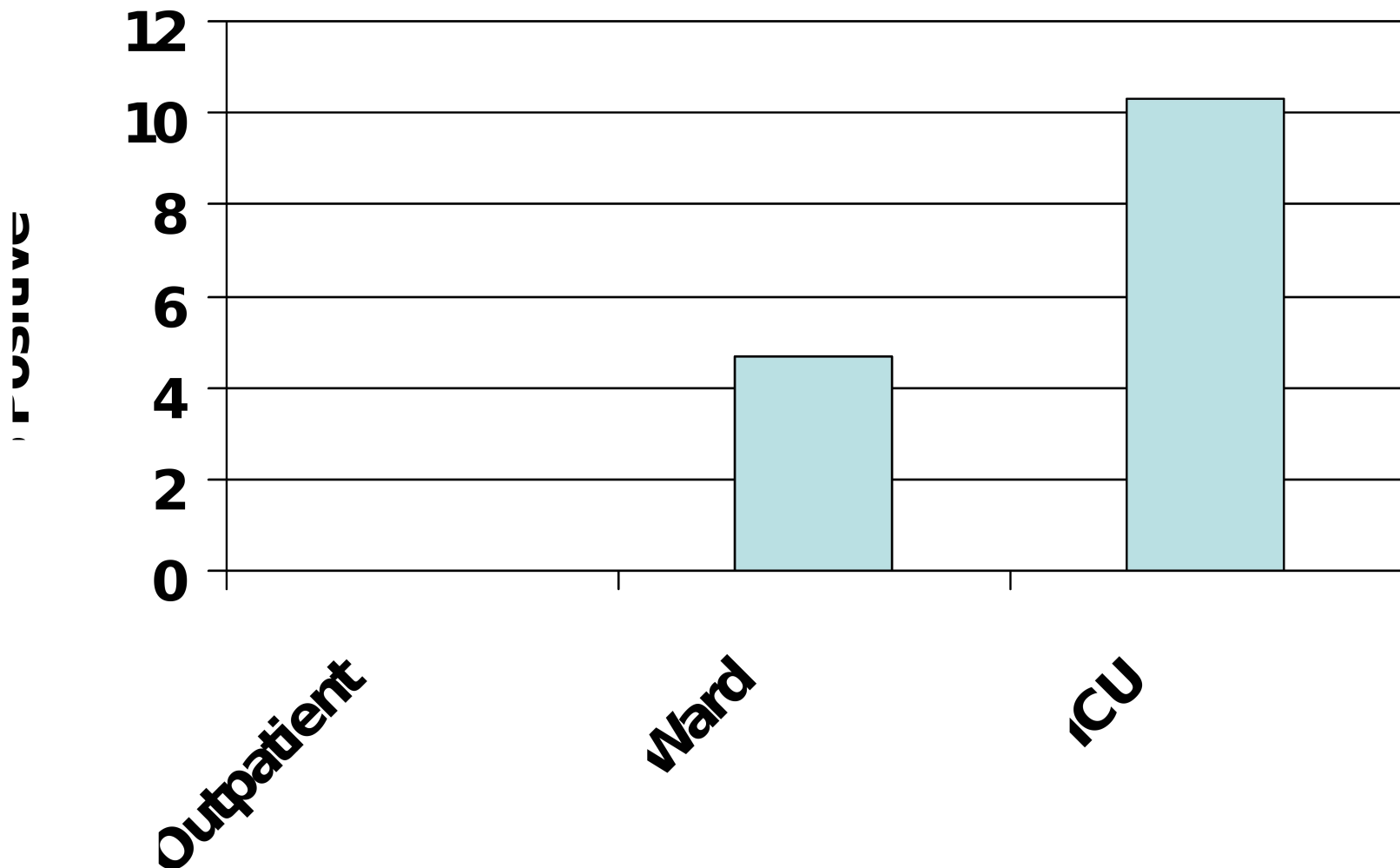


Acinetobacter Wound Cultures, LRMC, 2003-2005





Acinetobacter Colonizations by Patient Status on Arrival to LRMC



MTF locations where environmental sampling yielded *A. baumannii*

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Mosul

Tikrit

Balad

Baghdad

Iraq

Tigris River

Euphrates River

Syria

Iran

Jordan

Saudi Arabia

Kuwait

Kuwait City

Persian Gulf



Medical Facilities



0 37.5 75 150 225 300 Kilometers



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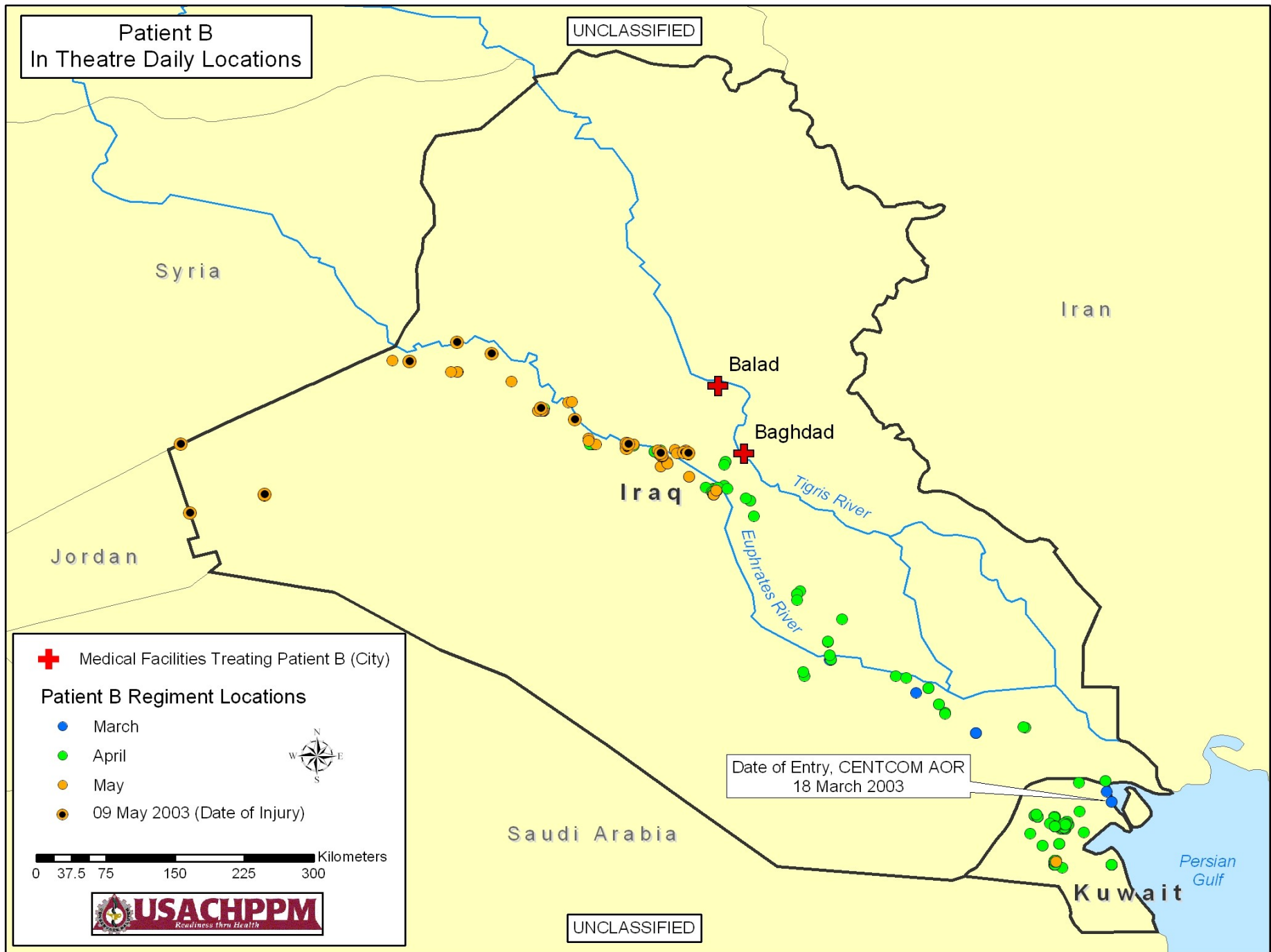


MEDEVAC Patient

- Isolate OIFC 138
 - Source: sputum
 - Date: 11 May 03
 - Location: LRMC
- Isolate OIFC 137
 - Source: catheter tip
 - Date: 23 May 03
 - Location: WRAMC
- 100% similar to isolate from MTF B Operating Room
- Evacuation Sequence
 - MTF E (Balad)
 - 10 May 03
 - MTF B (Dogwood)
 - 10 May 03
 - LRMC
 - 11 May 03
 - WRAMC
 - 22 May 03

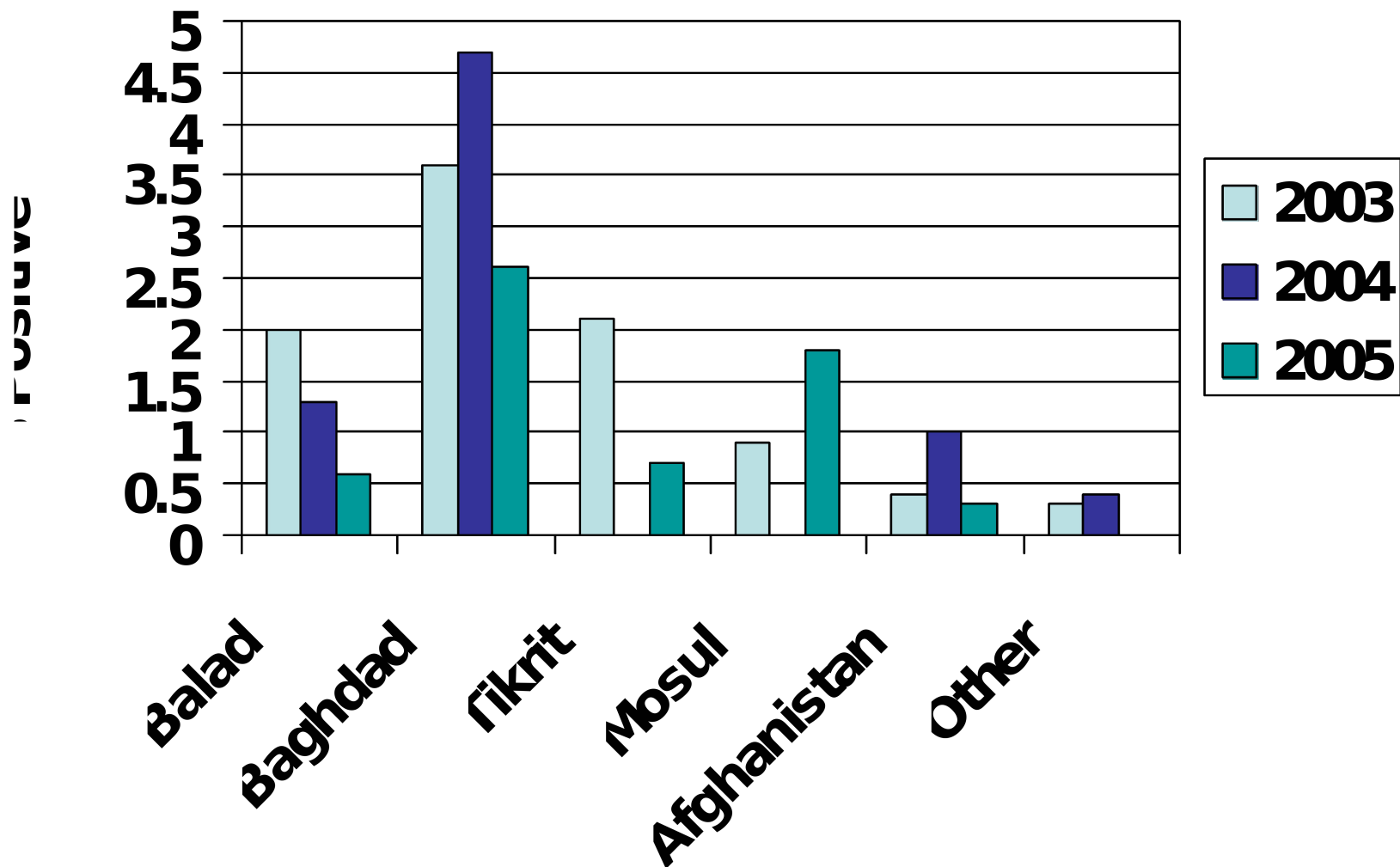
Patient B
In Theatre Daily Locations

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Acinetobacter Infection By Referring Level III Facility





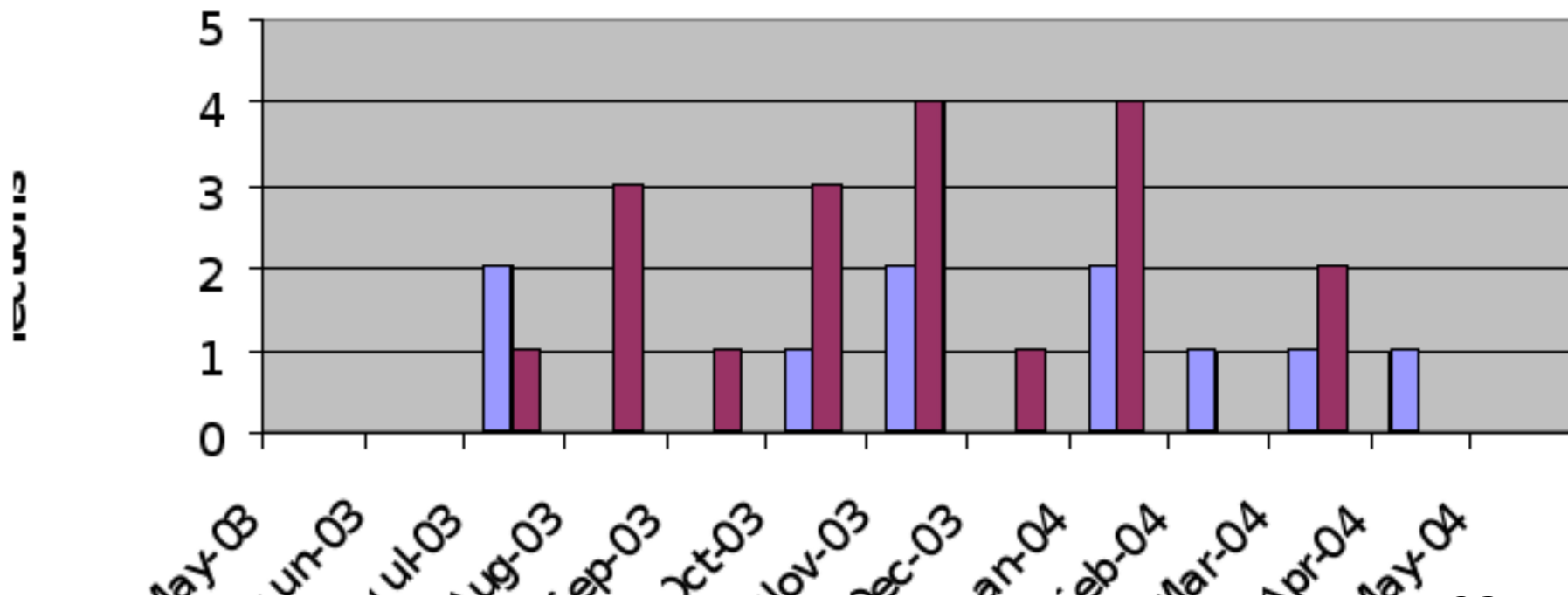
Nosocomial Patient

- LRMC
 - April 2004- 63yo woman admitted with COPD exacerbation
 - Admitted to med/surg ward (NOT ICU)
 - Developed Nosocomial Pneumonia due to MDR *Acinetobacter baumannii*.
 - Only *Acinetobacter* patient with bacteremia not from OIF
 - Died of Sepsis



Transmission at a Referral Hospital in Germany

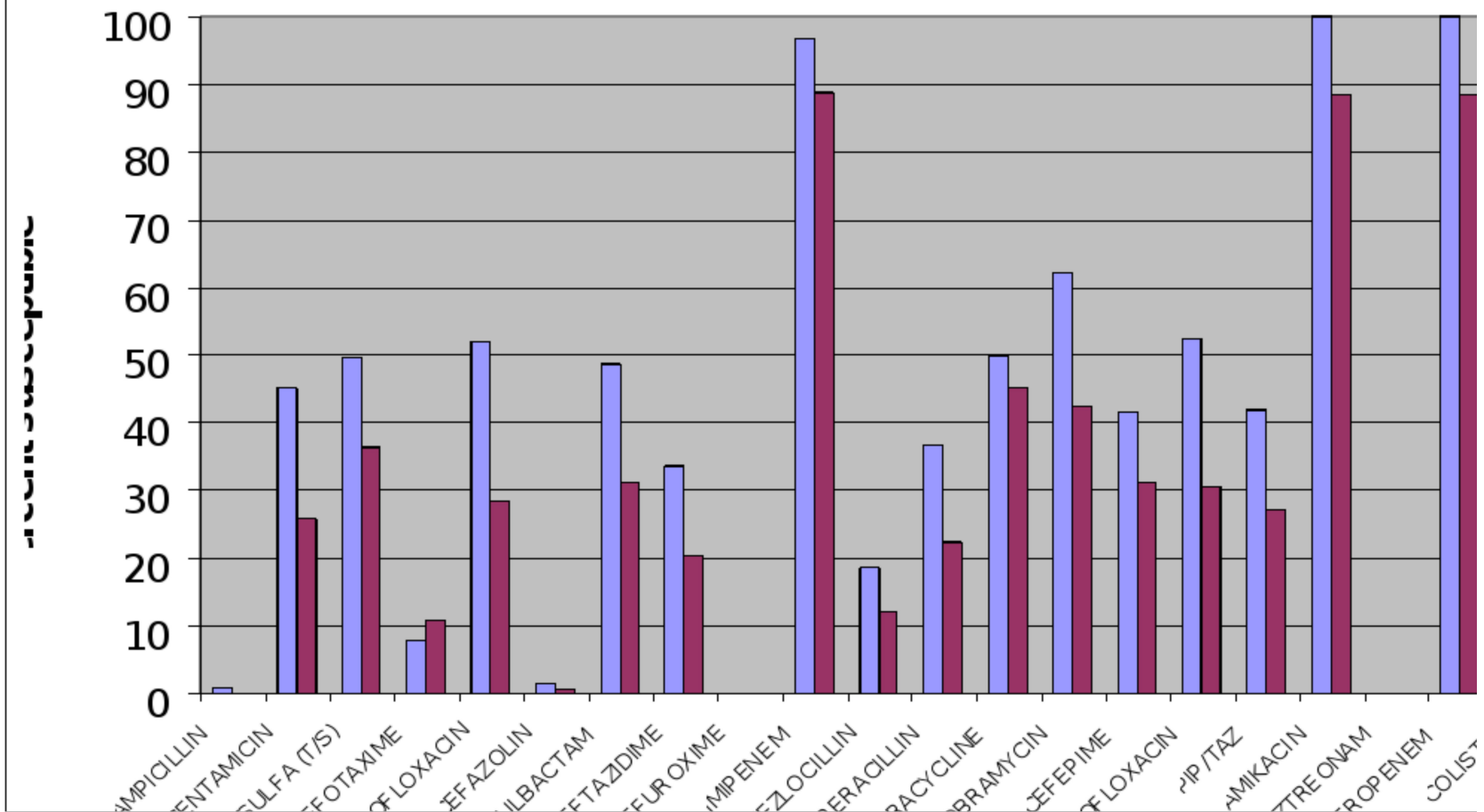
Epidemic curve of MDR Ab infection at University Saarland Hospital





Antibiotic Susceptibility

Antibiotic Susceptibility Comparison AB LRMC



Source: Greg Deye and Wade Aldous, LRMC



LRMC Admission Skin Isolates

Abx	2003 %S	2004 %S	p
Amp	2.083333	0	0.213
Gent	65.57377	32.76836	0.0001
T/S	64.58333	40.11299	0.0032
Cipro	68.85246	36.15819	0.0001
Amp/Sul	56.25	36	0.0475
Ceftaz	49.15254	25	0.001
Imi	97.91667	90.22989	0.1316
Pip	54.09836	28.24859	0.0005
TCN	70.96774	51.42857	0.0513
Tobr	73.77049	44	0.0001
Cefepime	51.6129	37.5	0.1646
Levoflox	65.71429	38.85714	0.0047
Amikacin	100	100	1
Meropenem	100	87.80488	0.5689

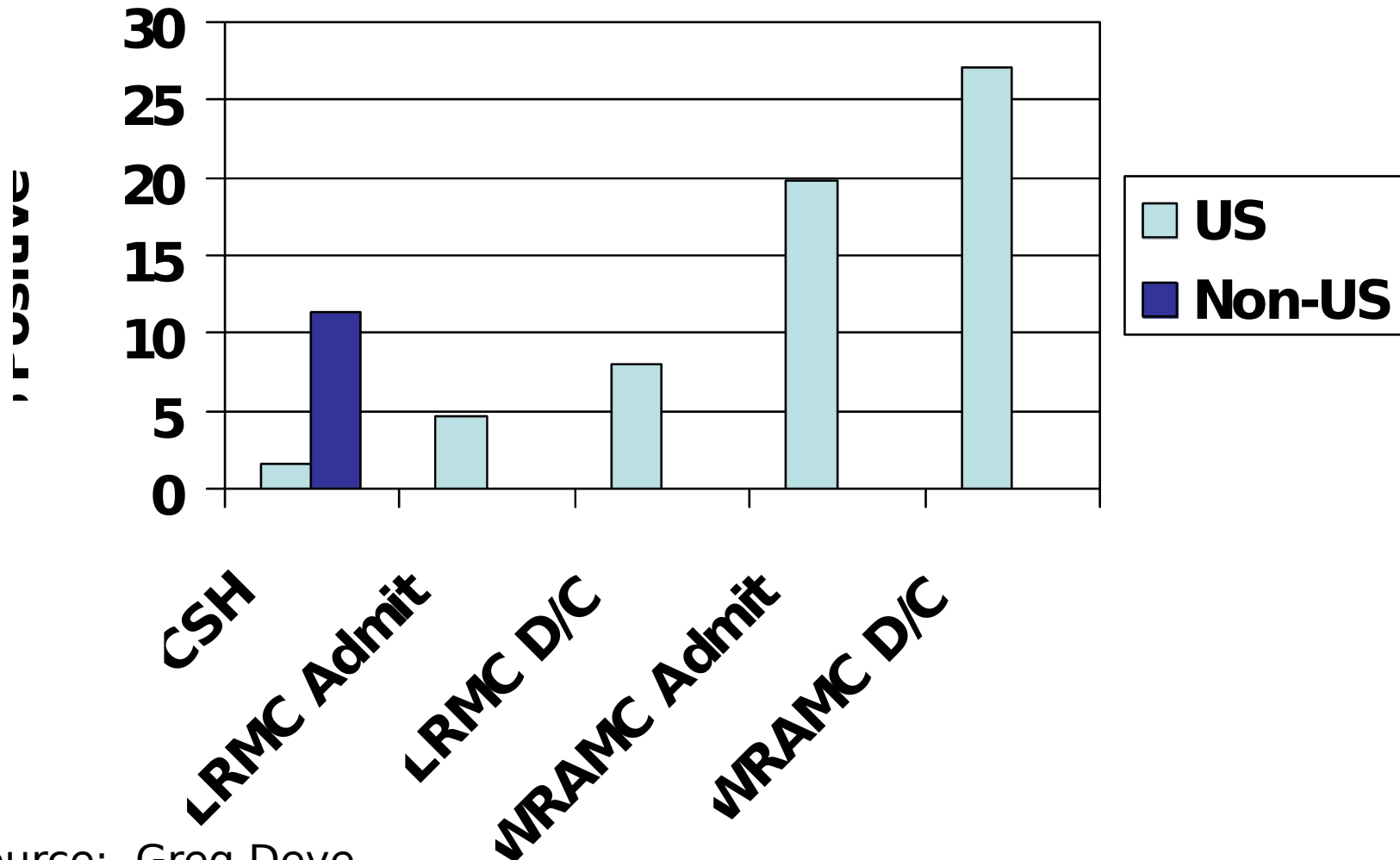


Case-Control Study of ICU Admissions With and Without Ab Bloodstream Infections

Variable	Odds Ratio	p-value
Transport>48 hrs	n/a	0.01
FST exposure	2.7	0.13
Baghdad	2.8	0.07
Antibiotic use	2.0	0.20
IED	0.8	0.80
Intubated	2.6	0.06
Blood products	5.0	0.00
Civilian	3.0	0.08



Acinetobacter Cultures at Selected Stages in the Military Healthcare System





Comparing “Success”

- Different culturing practices
 - Sites: Nares, wound, skin
 - Number of cultures
 - Access to culture capability
 - Culturing technique
 - Time to culture
 - Laboratory practices
- Uncertain relative contribution of nosocomial transmission, control measures throughout the MHS



Concordance Data

- Comparison of LRMC and WRAMC culture concordance from May to July 2005
- LRMC rates
 - Positive on admission = 4.0% (25/623)
 - Positive on discharge = 8.7% (13/149)
 - ICU patients on admission = 4.6% (8/176)
 - ICU patients on discharge = 8.3% (6/72)
- LRMC study group
 - Positive on admission = 3.6% (3/84)
 - Positive on discharge = 11.1% (4/36)
 - ICU admission = 4.2% (3/72)
 - ICU discharge = 10.7% (3/28)
- WRAMC study group, n=89
 - Positive on admission = 14.6% (13/89)
 - Axilla 10.1% (8/79)
 - Groin 12.3% (9/73)
 - ICU patients at LRMC = 66% (59/89)



Concordance Data, LRMC Admission vs. Discharge Skin Surveillance Cultures, May-July 2005

	D/C Cx +	D/C Cx -	Total
Admit Cx +	5	4	9
Admit Cx -	8	132	140
Total	13	136	149

McNemar's test,
 $p=0.25$

Kappa=0.41

LRMC Overall admission rate= 4.0%
This study = 6.0%



Concordance Data, LRMC Discharge vs. WRAMC Admission, Skin Surveillance Cultures, May-July 2005

	WRAMC Admit +	WRAMC Admit -	Total
LRMC D/C +	2	2	4
LRMC D/C -	2	30	32
Total	4	32	36

McNemar's test,
 $p=1.0$

Kappa=0.44

Note: 78% were ICU
patients at LRMC (28/36)



Concordance Data, WRAMC

Groin vs. Axilla Admission, Skin Surveillance Cultures, May-July 2005

	Groin Cx +	Groin Cx -	Total
Axilla Cx +	4	4	8
Axilla Cx -	5	50	55
Total	9	54	63



Infection Control Challenges

- Lack of hand hygiene products and sinks
 - 40-60% compliance with hand washing
- Shortage of gowns
- Repeated shortages of antibiotics
- Cohorting of patients to achieve infection control
- **Retroactive vs. proactive**
 - Wait for lab positives rather than up front isolation



Control Measures

- Surveillance
- Education
- Isolation
 - Not described in literature for GNRs (MRSA only)
 - No clear APIC, SHEA guidelines
 - MDR Pseudomonas clearly not transmitted person/person
 - MDR Acinetobacter-strong evidence of person-person and surfaces
- Decontamination
- Hand washing enforcement
- Barrier precautions
- Appropriate antibiotic use and guidelines
- Topical medications aimed at elimination?
- Cohort patients
- Shut down infected units



Campaign to Prevent Antimicrobial Resistance

Centers for Disease Control and Prevention

National Center for Infectious Diseases
Division of Healthcare Quality Promotion

Clinicians hold the solution!



12 Steps to Prevent Antimicrobial Resistance: Hospitalized Adults

Prevent Infection

1. Vaccinate
2. Get the catheters out

Diagnose and Treat Infection Effectively

3. Target the pathogen
4. Access the experts

Use Antimicrobials Wisely

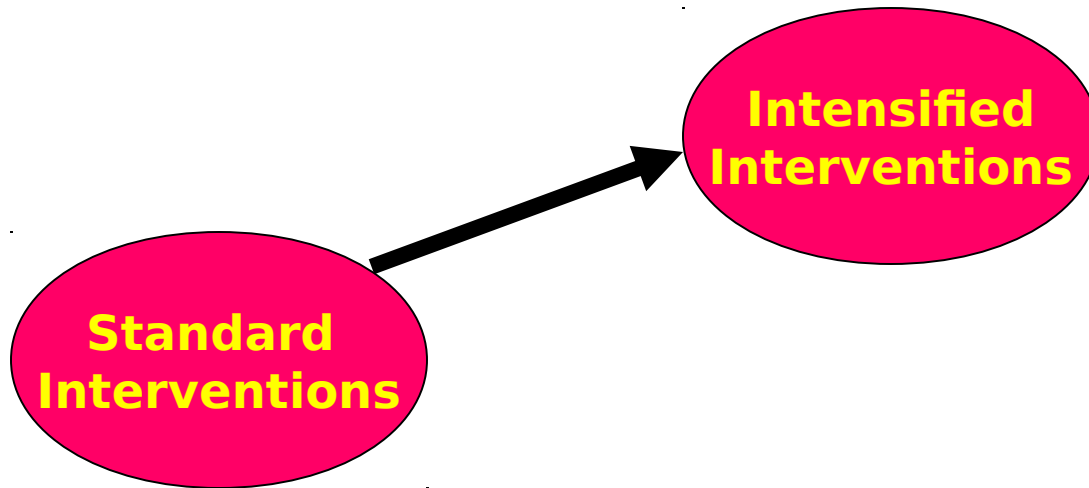
5. Practice antimicrobial control
6. Use local data
7. Treat infection, not contamination
8. Treat infection, not colonization
9. Know when to say “no” to vanco
10. Stop treatment when infection is cured or unlikely

Prevent Transmission

11. Isolate the pathogen
12. Break the chain of contagion



Draft HICPAC Isolation Guideline: Intensified MDRO Prevention Interventions



- **Evidence of ongoing transmission**
- **Prevalence of target MDRO has exceeding institutional goals**
- **First case of epidemiologically important MDRO**



Recommendations

- A dedicated IC practitioner at all level III and higher
- Standardized IC practices in place at all level III and higher facilities.
- Standardized surveillance with one technique (ax/groin) for all inpatients coming from theater for all MTFs, including level III. This implies microbiology capability able to support this surveillance.
- Standardized reporting requirement all level III and higher
- Improved documentation at all levels of care
- Inspection or reporting mechanisms where compliance with these measures will be verified



Conclusions

- *A. baumannii* is causing significant clinical disease
 - Once established, it is difficult to eradicate
 - Nosocomial spread and deaths have occurred
- Optimal Infection Control practices critical
- Field setting
 - Dedicated assets in field settings
 - Surveillance and lab capabilities in field setting
- MHS wide
 - Standardize surveillance for more meaningful comparisons